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(54) Title: COMBINATION OF AN IODOPROPYNYL DERIVATIVE WITH A KETONE ACID OR ITS SALT AND/OR WITH AN AROMATIC CARBOXYLIC ACID OR ITS SALT

(57) Abstract: Cyclic and acyclic ketone acids, such as dehydroacetic acid, and salts thereof as well as an aromatic carboxylic acid and salts thereof enhance the performance of iodine containing biocides as antimicrobial agents and preservatives. The present invention provides a composition comprising (a) an iodine containing biocide; and (b) (i) a ketone acid or salt thereof, (ii) an aromatic carboxylic acid or a salt thereof, or (iii) a mixture thereof. Preferably, the ketone acid is a cyclic ketone acid and the aromatic carboxylic acid is salicylic acid.

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COMBINATION OF AN IODOPROPYNYL DERIVATIVE WITH A KETONE ACID OR ITS SALT
AND/OR WITH AN AROMATIC CARBOXYLIC ACID OR ITS SALT

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15 This application claims the benefit of U.S. Provisional Application Serial
No. 60/273,079, filed March 1, 2001, which is hereby incorporated by reference.

Field of the Invention

20 This invention relates to antimicrobial compositions containing (a) an iodine
containing biocide, such as 3-iodo-2-propynyl butyl carbamate, and (b) (i) a cyclic or acyclic
ketone acid or a salt thereof, (ii) an aromatic carboxylic acid or a salt thereof, or (iii) a mixture
thereof.

Background of the Invention

25 Many iodine containing compounds, such as 3-iodo-2-propynyl butyl
carbamate (IPBC), are known to be effective as antimicrobial agents and preservatives.
However, IPBC and many other iodine containing compounds are expensive. As a result,

there is a continuing need for improved antimicrobial and preservative compositions which contain low concentrations of iodine containing biocides.

Summary of the Invention

5 Applicants have discovered that ketone acids, aromatic carboxylic acids, and salts thereof synergistically enhance the performance of iodine containing biocides. The present invention provides a composition comprising

(a) an iodine containing biocide; and

(b) (i) a ketone acid or salt thereof,

10 (ii) an aromatic carboxylic acid or a salt thereof, or

(iii) a mixture thereof.

Preferably, the ketone acid is a cyclic ketone acid. The aforementioned mixtures are synergistic.

Another embodiment of the present invention is a method for inhibiting the
15 growth of microorganisms on a substrate by applying an antimicrobial or preserving effective amount of the composition of the present invention.

Detailed Description of the Invention

The term "biocide", but is not limited to, bactericides, fungicides, pesticides
20 and agents which inhibit the growth of and/or destroy microorganisms and insects.

The present invention provides a composition comprising (a) an iodine containing compound; and (b) (i) a ketone acid or salt thereof, (ii) an aromatic carboxylic acid or a salt thereof, or (iii) a mixture thereof. Preferably, the ketone acid is a cyclic ketone acid. The ketone acid and aromatic carboxylic acid enhances the biocidal efficacy of the iodine
25 containing biocide. These compositions are useful as antimicrobial, fungicidal, and bactericidal agents and as preservatives in the papermaking, textile, agricultural, and coating industries and personal care, household, industrial, and institutional products, such as starches,

paints, adhesives, polyvinyl chloride and other plastics, and meltworking fluids. The preservative system may be incorporated into substrates susceptible to microbial growth. For example, the preservative system may be incorporated into or be a personal care product, such as a shampoo, conditioner, cream, lotion, cosmetic, and soap; a household product, such a
 5 fabric softener, laundry detergent, and hard surface cleaner; or an industrial product, such as paint, wood, textile, adhesive, sealant, leather, rope, paper pulp, plastic, fuel, oil, rubber working fluid, and metal working fluid.

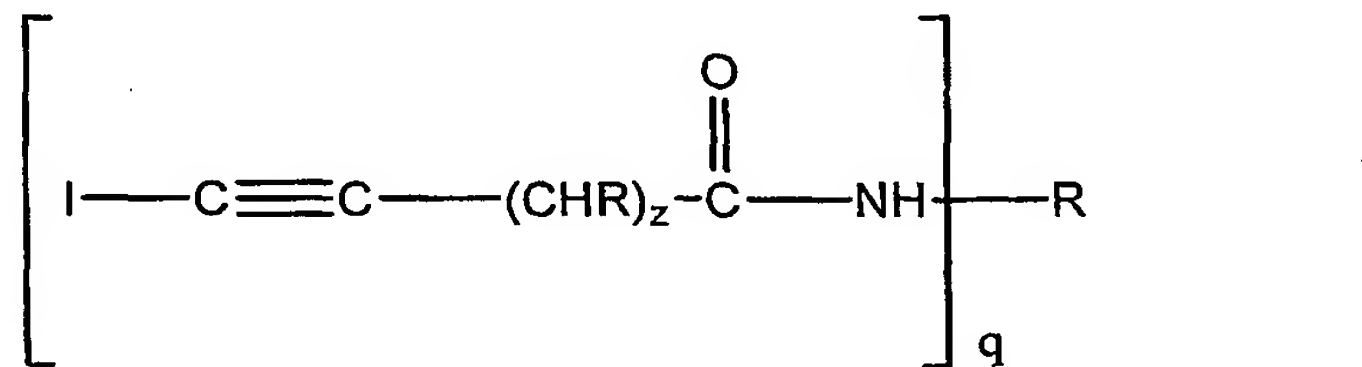
Examples of compounds which may be used as the iodine containing biocide component of the invention are fungicidally active iodoalkynyl derivatives. These include
 10 compounds derived from propyne or iodopropynyl alcohols, such as the esters, ethers, acetals, carbamates and carbonates and the iodopropynyl derivatives of pyrimidines, triazolinones, tetrazoles, triazinones, sulfamides, benzothiazoles, ammonium salts, carboxamides, hydroxamates, and ureas.

15 Iodine Containing Compounds

Preferred iodine containing biocides include, but not limited to, 3-iodo-2-propynyl derivatives such as 3-iodo-2-propynyl butyl carbamate, 3-iodo-2-propynyl succinate and p-chlorophenyl-3-iodopropynyl formal; iodo sulfone derivatives; and triiodoallyl alcohols.

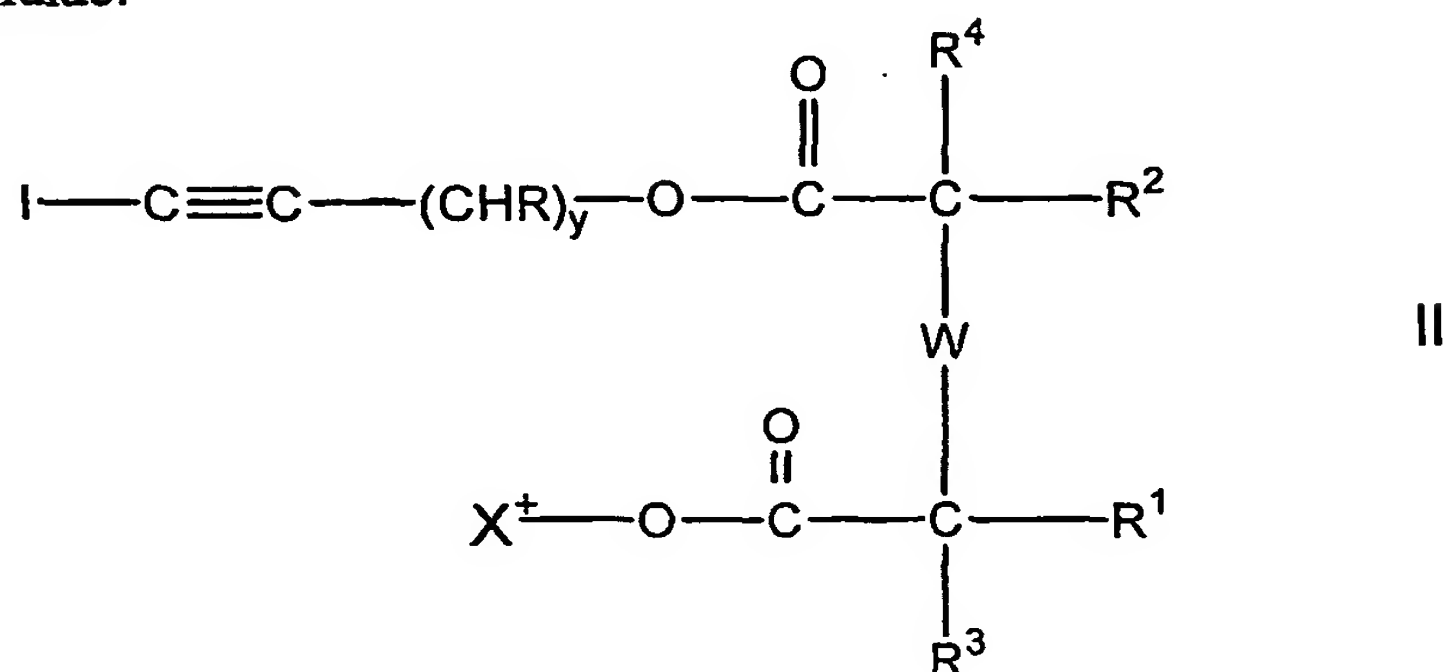
Preferably, iodopropynyl carbamate compounds have the formula:

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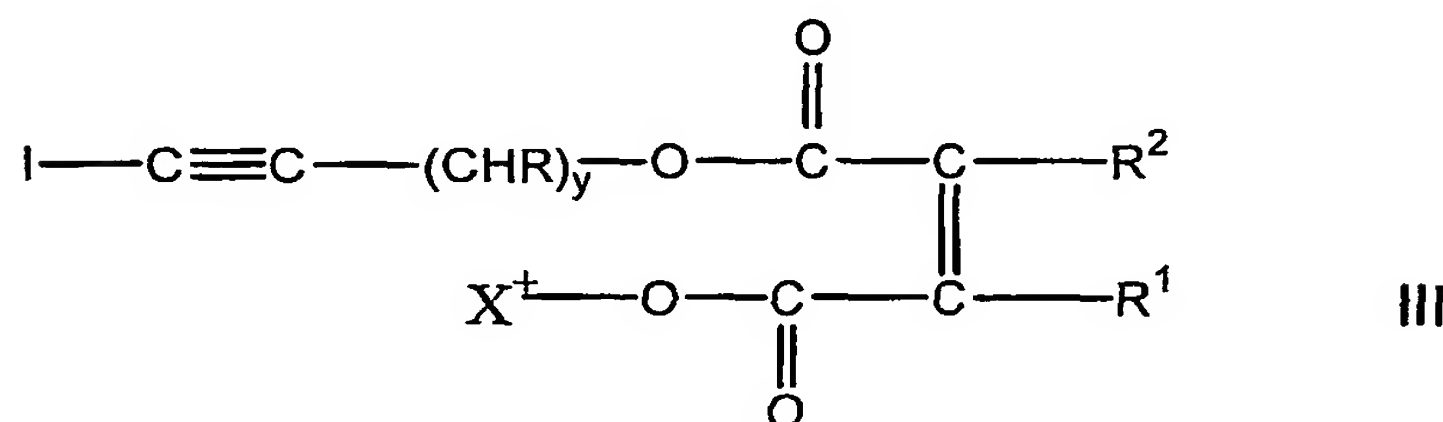


25 wherein R is selected from the group consisting of hydrogen, and substituted and unsubstituted alkyl, aryl, and alkylaryl groups having from 1 to 20 carbon atoms; and q and z independently are integers from 1 to 3.

Suitable iodopropynyl ester compounds include, but are not limited to, those of the following formulae:



and



15 wherein:

R¹ and R² are defined as R³ and R⁴ below or are joined to form a cycloalkyl, cycloalkenyl, aromatic or a heterocyclic ring containing an oxygen, nitrogen or sulfur atom or an alkoxy, amino, carboxyl, halo, hydroxyl, keto or a thiocarboxyl-substituted derivative thereof;

20 R^3 and R^4 are independently selected from

(A) hydrogen, alkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl, a heterocyclic ring containing an oxygen, nitrogen or sulfur atom, alkoxy, amino, carboxyl, halo, hydroxyl, keto or a thiocarboxyl; and

25 (B) substituted derivatives of the alkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl and the heterocyclic ring wherein the substitutions are alkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl,

alkoxy, amino, carboxyl, halo, hydroxyl, keto or a thiocarboxyl;

y is an integer from 0 to 16;

W is a single bond, oxygen, $-N(R^5)-$ or $-(CR^6R^7)_p-$;

5 R^5 is hydrogen, alkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl or a heterocyclic ring containing an oxygen, nitrogen or sulfur atom or a substituted derivative of alkyl, cycloalkyl, alkenyl, cycloalkenyl or aryl groups wherein the substitutions are alkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl, alkoxy, amino, carboxyl, halo, hydroxyl, keto, or a thiocarboxyl;

R^6 and R^7 are defined as R^3 and R^4 above;

10 p is an integer from 1 to 12; and

X is hydrogen or a salt-forming cation such as an alkali metal, an alkaline earth metal, ammonium, tertiary ammonium, a quaternary ammonium, a biguanide or a polybiguanide.

The above definition of R^5 includes, among other things, an aminoalkyl group.

15 The heterocyclic rings referred to in the above definitions may contain from 5 to 8 members, the alkyl or cycloalkyl groups from 1 to 18 atoms, the alkenyl or cycloalkenyl groups from 2 to 18 carbon atoms, and the aryl groups from 6 to 10 members.

In formula III, when R^1 and R^2 are hydrogen, the compound is a maleate. When R^1 and R^2 are joined together to form part of a six membered aromatic ring the compound is a phthalate. In formula II, when R^1 , R^2 , R^3 , and R^4 are hydrogen and W is a single bond, the compound is a succinate. When R^1 , R^2 , R^3 and R^4 are hydrogen and W is an oxygen, the compound is a diglycolate. Other compounds include the mono-iodopropynyl esters of anhydrides such as ethylenediamine tetraacetic dianhydride, 3,3-dimethylglutaric anhydride, S-acetylmercaptosuccinic anhydride, dichloromaleic anhydride, 2-dodecen-1-yl succinic anhydride and cis-5-norbornene-endo-2,3-dicarboxylic anhydride. Where hydrophilicity is desired, the sodium salts may be used because of their extremely high water solubility. Preferred carboxylic acid anhydrides include, but are not limited to, succinic,

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itaconic, phthalic, tetrachlorophthalic, and diglycolic anhydride. Examples of such compounds are described in U.S. Patent Nos. 4,844,891 and 5,073,570.

More preferably, the iodine containing biocide is 3-iodo-2-propynyl butyl carbamate (IPBC). The IPBC may be any grade of IPBC including, but not limited to, an essentially pure commercial grade IPBC in solid form and commercially available 6% and 10% grades in a solvent.

Another class of suitable iodopropynyl compounds are those having the formula:



where R^8 is benzyl or benzyl substituted with a methyl, methoxy, carboxyl, halogen or nitro group. A preferred compound is p-chlorophenyl-3-iodopropynyl formal.

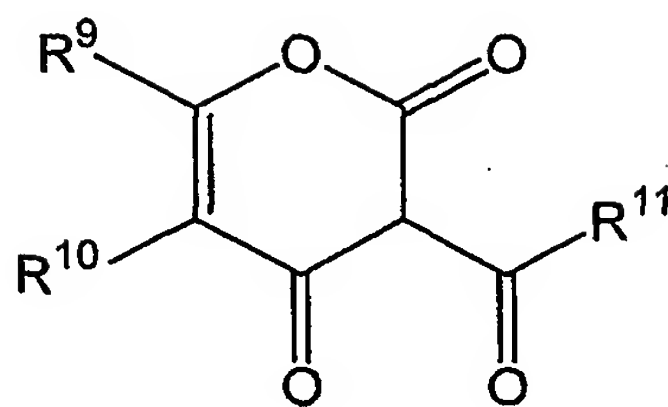
The iodine containing biocide may optionally be encapsulated, such as, for example, in cyclodextrin; calixarenes, such as 4-tert-butylcali[4]arene; liposomes; catezones; and amphiphilic betaine polymers. One example of an encapsulated iodine containing biocide is IPBC encapsulated in cyclodextrin, available as Troy Polyphase 604 from Troy Chemical Co. of East Hanover, NJ.

20 Ketone Acids

The ketone acid may be a cyclic or acyclic ketone acid. The term "cyclic ketone acid" as used herein includes compounds that have a ring containing a carbonyl group.

Suitable cyclic ketone acids include, but are not limited to, those having the formula

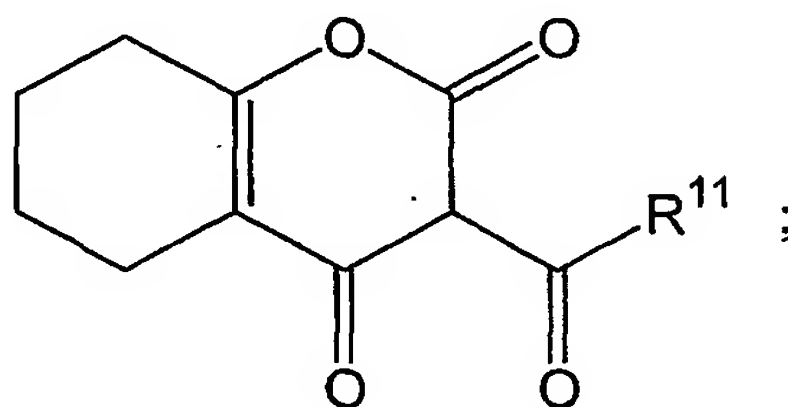
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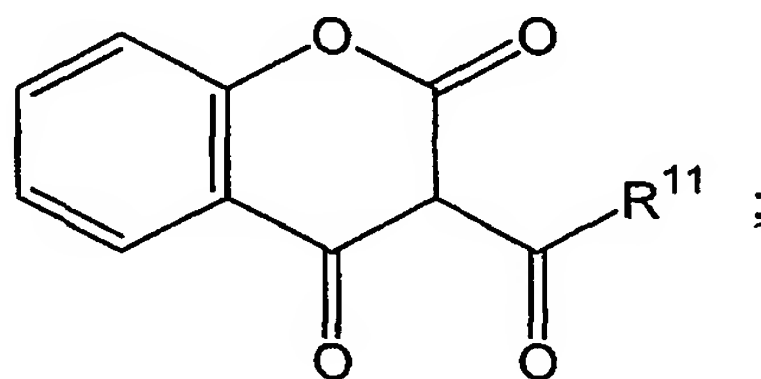
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and salts thereof, wherein R^9 , R^{10} , and R^{11} are independently C_1 - C_{10} alkyl, C_1 - C_{10} alkenyl, C_1 - C_{10} alkenyl, aryl, aryl substituted with halogen, or $(C_1$ - C_{10} alkyl)aryl. Preferably, R^9 , R^{10} , and R^{11} are independently C_1 - C_4 alkyl; or R^9 and R^{10} form a 5-12 member ring. Preferred cyclic ketone acids, include, but are not limited to, those having the formula

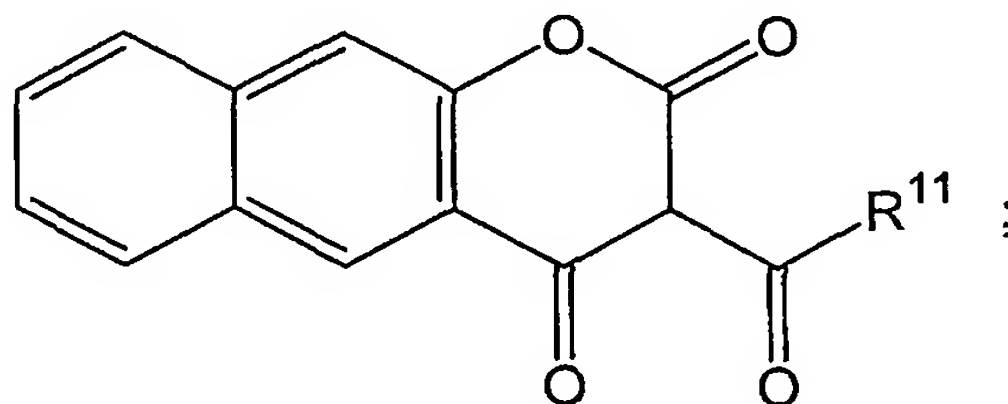
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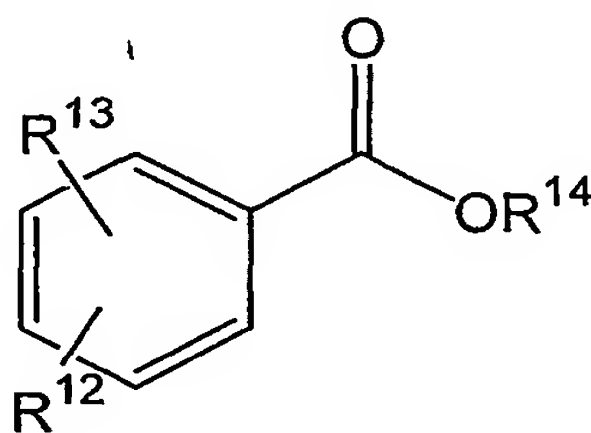
and salts thereof. A more preferred cyclic ketone acid is dehydroacetic acid and salts thereof (including hydrates thereof), such as sodium dehydroacetate (*e.g.* sodium dehydroacetate hydrate and sodium dehydroacetate monohydrate).

The cyclic ketone acid may optionally be encapsulated by any method in the art to increase its solubility in a desired solvent or formulation. For example, the cyclic ketone acid may be encapsulated in cyclodextrin; calixarenes, such as 4-tert-butylcali[4]arene; liposomes; catezones; and amphiphilic betaine polymers.

5 A preferred combination of cyclic ketone acid and iodine containing biocide is dehydroacetic acid and IPBC.

Aromatic Carboxylic Acids

Suitable aromatic carboxylic acids include, but are not limited to, benzoic acids, derivatives thereof, and salts thereof. According to one embodiment, the aromatic carboxylic acid has the formula



where R^{12} and R^{13} are independently H, -OH, or -OC(O)CH₃; and R^{14} is H, Na, K, Ca, or Mg. When R^{14} is Ca or Mg, the ratio of aromatic carboxylic acid to Ca or Mg may be 1:1 or 2:1.

20 For example, the aromatic carboxylic acid can be a hydroxy benzoic acid, derivative thereof, or salt thereof. A preferred hydroxy benzoic acid is salicylic acid and salts thereof. Suitable salts of salicylic acid include, but are not limited to, sodium salicylate.

A preferred combination of salicylic acid or salt thereof and iodine containing biocide is sodium salicylate and IPBC.

The composition may include a solvent, such as water and water miscible solvents, including, but not limited to, alcohols, glycols (e.g. glycerin, diglycerin, butylene

glycol, butoxydiglycol, propylene glycol, and dipropylene glycol), esters, ethers, polyethers, and any combination of any of the foregoing. For example, the solvent may comprise water and an alcohol, such as phenoxyethanol and/or benzyl alcohol.

Other adjuvants may be included in the composition as known to one of ordinary skill in the art. Suitable adjuvants include, but are not limited to, preservatives; solubilizing agents; chelating agents, such as ethylenediaminetetraacetic acid (EDTA) and salts thereof and zeolites; surfactants, such as cationic, anionic, nonionic, and amphoteric surfactants; antioxidants, such as butylated hydroxyanisole (BHA) and butylhydroxytoluene (BHT); amine oxides; tertiary amines; hydrotropes; zinc compounds; hydrotropes; fluoride compounds; magnesium salts; calcium salts; carboxylic acids; phosphates; phosphonates; formaldehyde donors; glycereth-7; myristyl myristate; glutaraldehydes; biguanides; natural products, such as usnic acid and tea tree oils; and any combination of any of the foregoing.

Suitable preservatives include, but are not limited to, quaternary ammonium chlorides, such as benzethonium chloride (available as Hyamine[®] 1622 from Lonza Inc. of Fair Lawn, NJ) and benzalkonium chlorides (available as Barquat[®] MB-50 and MB-80 from Lonza Inc. of Fair Lawn, NJ); hydantoins, such as dimethylhydantoin and halogenated hydantoins; isothiazolinones; parabens, such as methylparaben, ethylparaben, and propylparaben; chloroxylenol; chlorhexidine; phenoxyethanol; benzyl alcohol; phenethyl alcohol; benzoic acid and salts thereof; chlorobutanol; sorbic acid and salts thereof; triclosan; triclocarban; and any combination of any of the foregoing.

Typically, the composition is an aqueous or oil based system and is not an emulsion. For compositions which are oil based, the iodine containing biocide is preferably not encapsulated and the ketone acid is preferably not a hydrate. A suitable solvent for an oil based system is phenoxyethanol. For compositions which are water based, the iodine containing biocide is preferably encapsulated to enhance its water solubility and the ketone acid is preferably a hydrate.

The composition can be a liquid or a solid.

The weight ratio of (1) ketone acid, aromatic carboxylic acid, or salts thereof or mixtures thereof to (2) iodine containing biocide broadly ranges from about 0.0006:1 to about 1990:1 and preferably ranges from about 0.0063:1 to about 1400:1. According to another embodiment, the molar ratio ranges from 0.063:1 to about 140:1 or from about 0.63:1 to about 14:1.

To prepare a formulation containing the composition of the present invention, a concentrate is generally first prepared. Table A illustrates the components and the ranges of components present in a typical concentrate (based upon 100% total weight of concentrate).

Table A

Ranges	Iodine Containing Biocide	Ketone Acid, Aromatic Carboxylic Acid, Salts Thereof, or Mixtures Thereof
Broad	from about 0.05 to about 80%	from about 0.05 to about 99.5%
Preferred	from about 0.5 to about 30%	from about 0.50 to about 70%
More Preferred	from about 1 to about 15%	from about 5 to about 40%

Before use, the concentrate is diluted, preferably with the same solvent as was used in the concentrate. Use dilutions of the composition typically comprise a biocidally, fungicidally, or bactericidally effective amount of (1) the iodine containing biocide (i.e., component (a)) and/or (2) the mixture of components (a) and (b) (where component (b) is the ketone acid, aromatic carboxylic acid, or salt thereof, or a mixture thereof). The use dilutions also typically comprise a biocidal, fungicidal, or bactericidal enhancing (or potentiating) effective amount of the ketone acid or salt thereof, aromatic carboxylic acid or salt thereof, or mixture thereof (i.e., component (b)). Generally, use dilutions contain from about 0.0001%, 0.01%, or 0.1% to about 2% by weight of the concentrate. According to one

preferred embodiment, use dilutions contain from about 0.1 to about 0.5% or 1% by weight of the concentrate.

Table B illustrates the components and generally the ranges of components present in the use dilution (based upon 100% total weight of use dilution).

Table B

Ranges	Iodine Containing Biocide	Ketone Acid, Aromatic Carboxylic Acid, Salts Thereof, or Mixtures Thereof
Broad	from about 0.00005 to about 0.40%	from about 0.00005 to about 0.4975%
Preferred	from about 0.0005 to about 0.15%	from about 0.0005 to about 0.35%
More Preferred	from about 0.001 to about 0.075%	from about 0.005 to about 0.2%

Another embodiment of the present invention is a method for inhibiting the growth of microorganisms, bacteria (e.g., *S. aureus* (ATCC # 6538), *P. aeruginosa* (ATCC # 9027), and *E. coli* (ATCC # 8739)), and/or fungi (e.g., *Candida albicans* and *Aspergillus niger*) on a substrate by applying an antimicrobial, bactericidal, or fungicidal effective amount of the composition of the present invention to the substrate. The composition may be applied to the substrate by any method known to one of ordinary skill in the art including, but not limited to, brushing, dipping, soaking, vacuum impregnation, and pressure treatment.

The composition of the present invention may be prepared by mixing the ketone acid or salt thereof and/or the aromatic carboxylic acid or salt thereof, the iodine containing biocide, solvents, and adjuvants. The mixture may be heated and/or stirred to expedite mixing.

Description of the Preferred Embodiments

The following examples illustrate the invention without limitation. All parts and percentages are given by weight unless otherwise indicated.

Example 1

Each anionic shampoo sample in Table 1 below was tested as follows. A standardized mixed bacterial solution was prepared according to the following procedure. 3 agar stabs of *S. Aureus* (ATCC # 6538), *P. aeruginosa* (ATCC # 9027), and *E. Coli* (ATCC # 8739) were separately incubated at about 35°C for about 24 hours. Each stab was then washed with 3 mL of sterile 0.85% saline solution. The washes of the 3 stabs were pooled together to form an organism mixture. The absorbance of the organism mixture at 530 nm was adjusted to about 1.00 by adding saline. The spectrometer was calibrated with a saline blank. A 5 mL aliquot of the organism mixture was mixed together to produce the standardized mixed bacterial solution. Then, 40 g of each shampoo sample was inoculated with 0.2 mL of the standardized mixed bacterial solution and mixed. 1 g of the mixture was added to a sterile 20 x 150 mm screw cap test tube.

9 mL of sterile D/E neutralizer broth was added to the test tube and mixed to form a 10^{-1} dilution. Serial dilutions were prepared through to a 10^{-6} dilution with phosphate buffered water. The serial dilutions were plated onto Tryptic Soy Agar and incubated for 2 days at about 35°C. Bacteria counts were performed after 0 and 14 days. The results are shown in Table 1.

The anionic protein shampoo composition was comprised of 35% by weight of sodium lauryl ether sulfate; 25% by weight of triethanolamine lauryl sulfate; 3% by weight coconut diethanolamide (cocamide DEA); 1% by weight of hydrolyzed collagen, available as Polypro 5000™ from Hormel Foods of Austin, MN; and 36% by weight of deionized water.

The sodium dehydroacetate monohydrate, sodium salicylate, and IPBC (Glycasil® 2000) shampoo samples were prepared by mixing the appropriate amounts of the

preservatives and the aforementioned protein shampoo composition and heating the mixture to about 50°C for about 15 minutes.

Table 1

Shampoo	<i>S. aureus</i> , <i>P. aeruginosa</i> , and <i>E. coli</i> (cfu/g)	
	Day 0	Day 14
Unpreserved Protein Shampoo Composition	3.0×10^6	3.0×10^7
0.5% Sodium Salicylate ¹ and 0.5% Glycakil [®] 2000 ² *	3.0×10^6	< 10
0.10% Sodium Dehydroacetate Monohydrate ³ , 0.10% Sodium Salicylate ¹ , and 0.50% Glycakil [®] 2000 ² *	3.0×10^6	< 10
0.25% Sodium Dehydroacetate Monohydrate ³ , 0.25% Sodium Salicylate ¹ , and 0.25% Glycakil [®] 2000 ² *	3.0×10^6	< 10
> 0.5% Sodium Dehydroacetate Monohydrate ³ *	3.0×10^6	4.0×10^3
> 1.0% Sodium Salicylate ¹ *	3.0×10^6	5.0×10^2
> 1.0% Glycakil [®] 2000 ² *	3.0×10^6	1.0×10^7

All percentages in Table 1 are in percent by weight based upon 100% by weight of total shampoo.

¹ - Sodium dehydroacetate monohydrate is available from Lonza Inc. of Fair Lawn, NJ.

² - Glycakil[®] 2000 is iodopropynyl butylcarbamate and is available from Lonza Inc. of Fair Lawn, NJ.

³ - Sodium salicylate is available from Sigma Chemical Co. of St. Louis, MO.

* - Below the specified concentration of preservative, the shampoo contained ≥ 10 cfu/g after 14 days.

Synergism for the sodium salicylate/Glycakil® 2000 solution in Table 1 against *S. aureus*, *P. aeruginosa*, and *E. coli* was calculated by the method described in C.E. Kull *et al.*, "Mixtures of Quaternary Ammonium Compounds and Long-chain Fatty Acids as Antifungal Agents", *Applied Microbiology*, 9:538-541 (1961). The synergism value ($Q_A/Q_a + Q_B/Q_b$) was determined. Q_A is the concentration of sodium salicylate (in percent by weight) in the mixture, which yielded 100% retardation of the bacteria, *i.e.*, resulted in a plate count of < 10 cfu/g after 14 days. Q_a is the concentration of sodium salicylate alone (in percent by weight) required to yield 100% retardation of the bacteria. Q_B is the concentration of Glycakil® 2000 (in percent by weight) in the mixture, which yielded 100% retardation of the bacteria. Q_b is the concentration of Glycakil® 2000 alone (in percent by weight) required to yield 100% retardation of the bacteria.

When the value of ($Q_A/Q_a + Q_B/Q_b$) is less than one, the mixture is synergistic. Values for ($Q_A/Q_a + Q_B/Q_b$) of 1 and greater than 1, represent an additive effect and an antagonistic effect, respectively.

The results are shown in Table 2 below.

Table 2

Preservative Mixture	Q_A	Q_B	Q_a	Q_b	$Q_A/Q_a + Q_B/Q_b$
0.5% Sodium Salicylate and 0.5% Glycakil® 2000	0.5%	0.5%	$>1.00\%$	$>1.00\%$	< 1

The synergism for the sodium dehydroacetate monohydrate/sodium salicylate/Glycakil® 2000 solutions in Table 1 was also calculated by this method. Q_A , Q_B , Q_a , and Q_b are defined as above. Q_C is the concentration of sodium dehydroacetate monohydrate (in percent by weight) in the mixture, which yielded 100% retardation of the bacteria. Q_c is the concentration of sodium dehydroacetate monohydrate alone (in percent by weight) required to yield 100% retardation of the bacteria. When the value of $(Q_A/Q_a + Q_B/Q_b + Q_C/Q_c)$ is less than one, the mixture is synergistic. Values for $(Q_A/Q_a + Q_B/Q_b + Q_C/Q_c)$ of 1 and greater than 1, represent an additive effect and an antagonistic effect, respectively.

The results are shown in Table 3.

Table 3

Coefficient	0.10% Sodium Salicylate, 0.50% Glycakil® 2000, and 0.10% Sodium Dehydroacetate Monohydrate	0.25% Sodium Salicylate, 0.25% Glycakil® 2000, and 0.25% Sodium Dehydroacetate Monohydrate
Q_A	0.10%	0.25%
Q_B	0.50%	0.25%
Q_C	0.10%	0.25%
Q_a	> 1.00%	> 1.00%
Q_b	> 1.00%	> 1.00%
Q_c	> 0.50%	> 0.50%
$Q_A/Q_a + Q_B/Q_b + Q_C/Q_c$	< 0.8	< 1

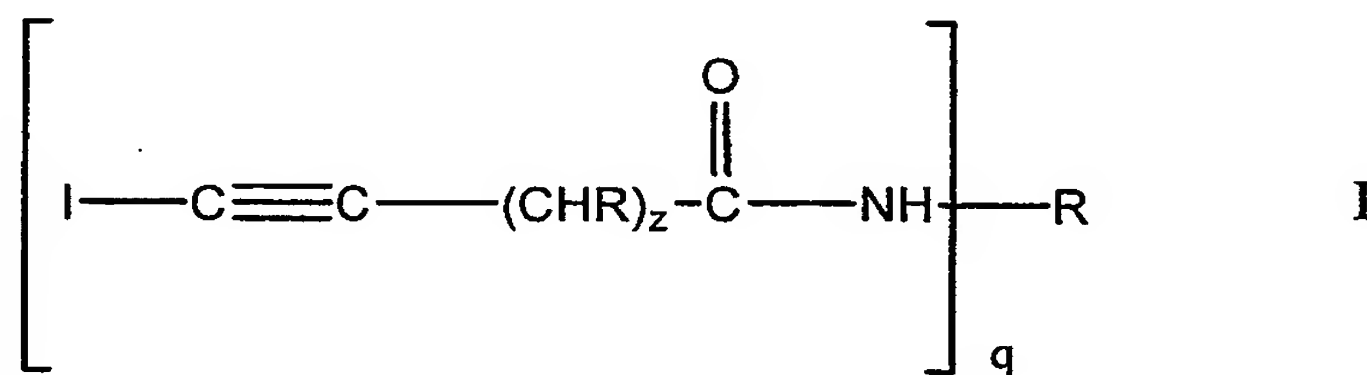
All patents, applications, articles, publications, and test methods mentioned above are hereby incorporated by reference.

Many variations of the present invention will suggest themselves to those skilled in the art in light of the above detailed description. Such obvious variations are within the full intended scope of the appended claims.

IN THE CLAIMS:

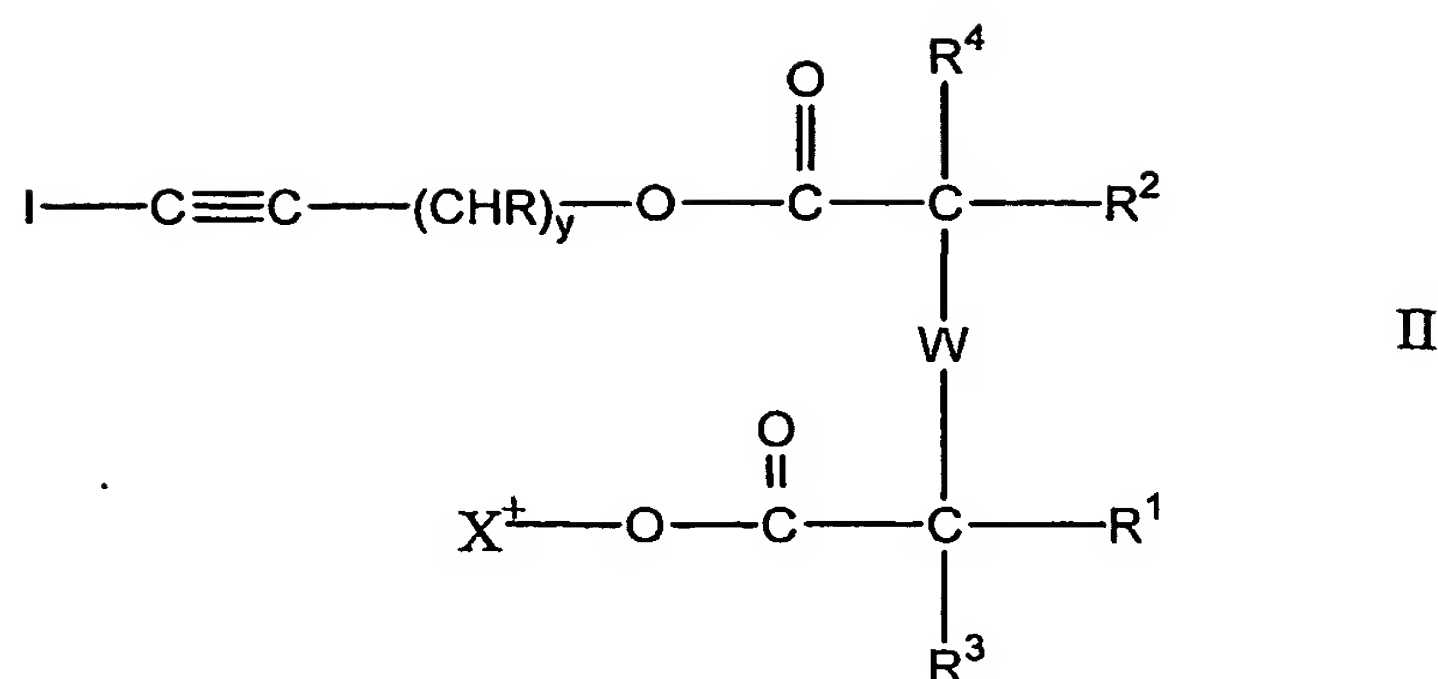
1. A composition comprising
(a) an iodine containing biocide; and
(b) (i) a ketone acid or salt thereof, (ii) an aromatic carboxylic acid or a salt thereof, or (iii) a mixture thereof.

2. The composition of claim 1, wherein the iodine containing biocide is an iodopropynyl derivative of the formula

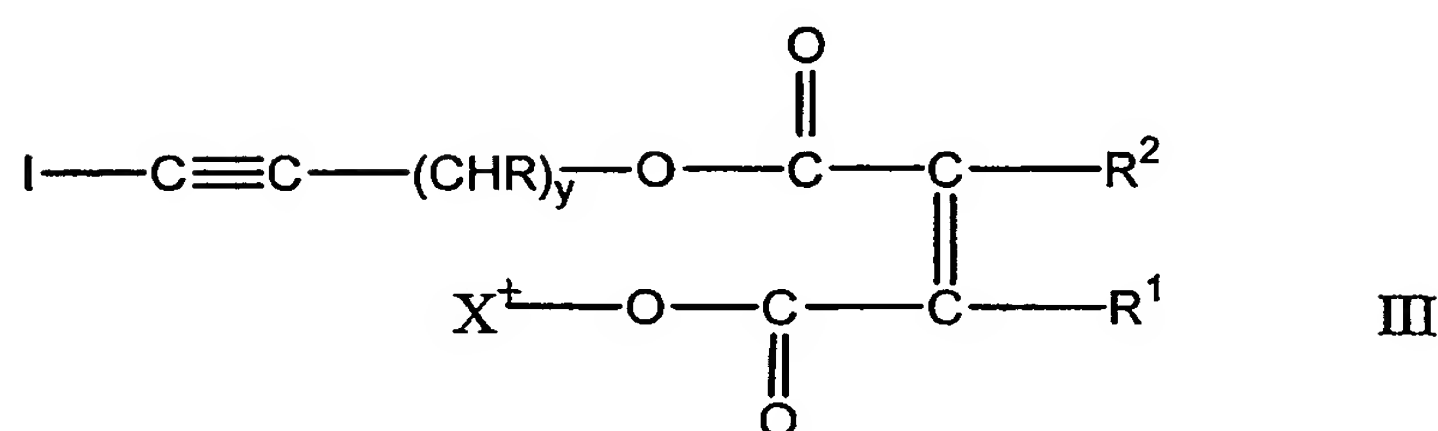


wherein R is hydrogen or a substituted or unsubstituted alkyl, aryl, or alkylaryl group having from 1 to 20 carbon atoms; and z and q are independent integers from 1 to 3.

3. The composition of claim 1, wherein the iodine containing biocide is an iodopropynyl derivative of the formula



or



wherein

R^1 and R^2 are defined as R^3 and R^4 below or are joined to form a cycloalkyl, cycloalkenyl, aromatic or a heterocyclic ring containing an oxygen, nitrogen or sulfur atom or an alkoxy, amino, carboxyl, halo, hydroxyl, keto or a thiocarboxyl-substituted derivative thereof;

R^3 and R^4 are independently selected from (A) hydrogen, alkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl, a heterocyclic ring containing an oxygen, nitrogen or sulfur atom, alkoxy, amino, carboxyl, halo, hydroxyl, keto or a thiocarboxyl and (B) substituted derivatives of the alkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl and the heterocyclic ring wherein the substitutions are alkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl, alkoxy, amino, carboxyl, halo, hydroxyl, keto or a thiocarboxyl; y is 0 to 16; W may be a single bond, oxygen, NR^5 , or $(\text{CR}^6\text{R}^7)_p$, wherein R^5 is hydrogen, alkyl, aminoalkyl, cycloalkyl, alkenyl, cycloalkenyl, aryl or a heterocyclic ring containing an oxygen, nitrogen or sulfur atom or a substituted derivative of alkyl, cycloalkyl, alkenyl, cycloalkenyl or aryl groups wherein the substitutions are alkyl,

cycloalkyl, alkenyl, cycloalkenyl, aryl, alkoxy, amino, carboxyl, halo, hydroxyl, keto, or a thiocarboxyl wherein R^6 and R^7 are defined as R^3 and R^4 above and p is an integer from 1 to 12; and wherein the heterocyclic rings comprise from 5 to 8 members, the alkyl or cycloalkyl groups from 1 to 18 atoms, the alkenyl or cycloalkenyl groups from 2 to 18 carbon atoms, and the aryl groups from 6 to 10 members.

4. The composition of claim 2, wherein the iodopropynyl derivative is a 3-iodo-2-propynyl derivative.

5. The composition of claim 4, wherein the 3-iodo-2-propynyl derivative is 3-iodo-2-propynyl butyl carbamate, 3-iodo-2-propynyl succinate or p-chlorophenyl-3-iodopropynyl formal.

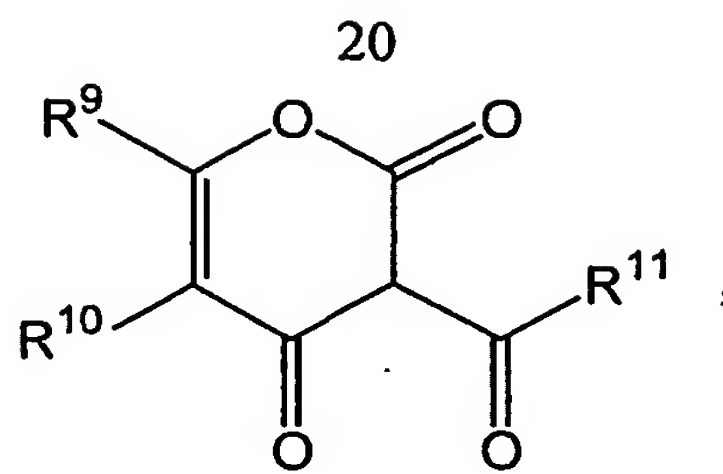
6. The composition of claim 5, the 3-iodo-2-propynyl derivative is 3-iodo-2-propynyl butyl carbamate.

7. The composition of claim 1, wherein the iodine containing biocide is encapsulated in cyclodextrin.

8. The composition of claim 10, wherein the 3-iodo-2-propynyl butyl carbamate is encapsulated in cyclodextrin.

9. The composition of claim 1, wherein the ketone acid is a cyclic ketone acid or a salt thereof.

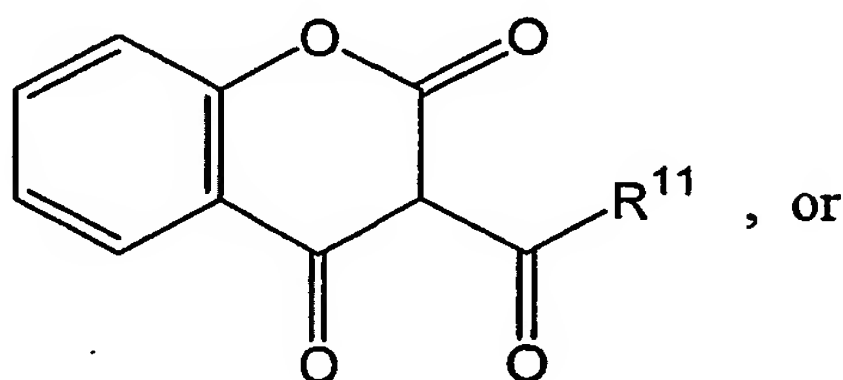
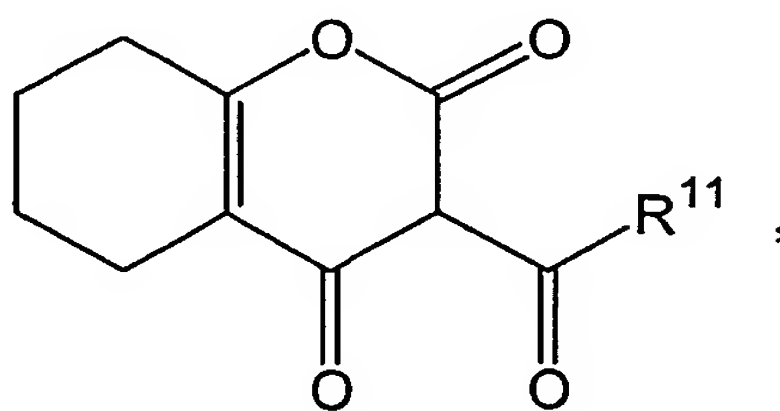
10. The composition of claim 9, wherein the cyclic ketone acid has the formula

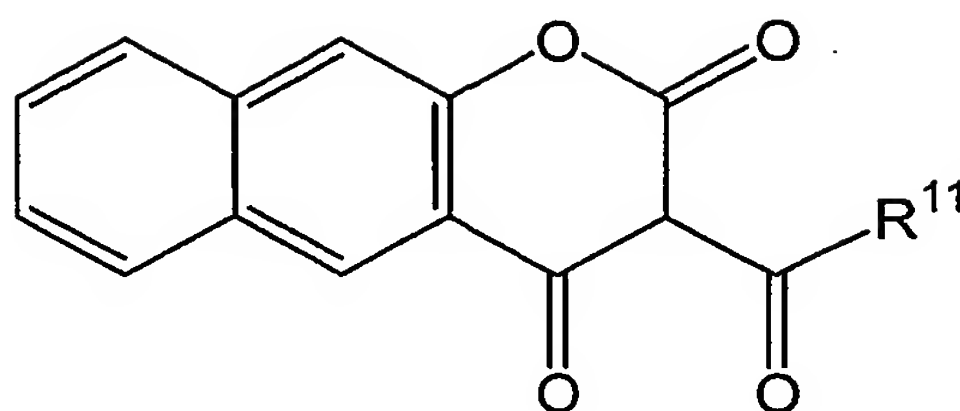


wherein R^9 , R^{10} , and R^{11} are independently C_1 - C_{10} alkyl, C_1 - C_{10} alkenyl, C_1 - C_{10} alkynyl, aryl, aryl substituted with halogen, or (C_1 - C_{10} alkyl)aryl.

11. The composition of claim 10, wherein R^9 , R^{10} , and R^{11} are independently C_1 - C_4 alkyl; or R^9 and R^{10} form a 5-12 member ring.

12. The composition of claim 10, wherein the cyclic ketone acid has the formula





13. The composition of claim 9, wherein the ketone acid is dehydroacetic acid or a salt thereof.

14. The composition of claim 1, wherein the ketone acid is sodium dehydroacetate.

15. The composition of claim 1, wherein the ketone acid is encapsulated.

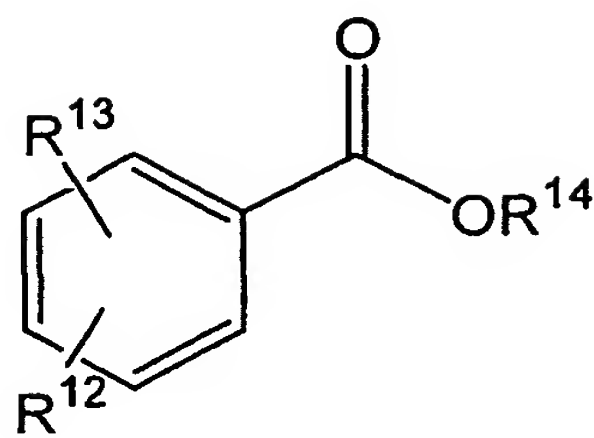
16. The composition of claim 13, wherein the dehydroacetic acid or salt thereof is encapsulated in cyclodextrin.

17. The composition of claim 1, wherein the iodine containing biocide is iodopropynyl butylcarbamate and the ketone acid is dehydroacetic acid or a salt thereof.

18. The composition of claim 1, wherein the aromatic carboxylic acid is benzoic acid, derivative thereof, or salt thereof.

19. The composition of claim 1, wherein the aromatic carboxylic acid has the formula

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wherein R¹² and R¹³ are independently H, -OH, or -OC(O)CH₃; and R¹⁴ is H, Na, K, Ca, or Mg.

20. The composition of claim 1, wherein the aromatic carboxylic acid is a hydroxy benzoic acid, derivative thereof, or salt thereof.

21. The composition of claim 20, wherein the hydroxy benzoic acid is salicylic acid or a salt thereof.

22. The composition of claim 21, wherein the salt of salicylic acid is sodium salicylate.

23. The composition of claim 1, wherein the iodine containing biocide is iodopropynyl butylcarbamate and the aromatic carboxylic acid is sodium salicylate.

24. The composition of claim 1, further comprising a solvent.

25. The composition of claim 24, wherein the solvent is water, an alcohol, a glycol, an ester, an ether, a polyether or any combination of any of the foregoing.

26. The composition of claim 24, wherein the solvent comprises water and alcohol.

1 27. The composition of claim 24, wherein the alcohol is phenoxyethanol.

1 28. The composition of claim 1, wherein the composition comprises a biocidally
2 effective amount of the iodine containing biocide.

1 29. The composition of claim 1, wherein the composition comprises a fungicidally
2 effective amount of the iodine containing biocide.

1 30. The composition of claim 1, wherein the weight ratio of the ketone acid to the
2 iodine containing biocide ranges from about 0.0006:1 to about 1990:1.

1 31. The composition of claim 30, wherein the weight ratio of the ketone acid to
2 the iodine containing biocide ranges from about 0.0063:1 to about 1400:1.

1 32. The composition of claim 1, wherein said composition is a use dilution
2 comprising from about 0.00005 to about 0.4975% by weight of ketone acid and from about
3 0.00005 to about 0.40% by weight of iodine containing biocide, based upon 100% weight of
4 total composition.

1 33. The composition of claim 32, wherein said composition is a use dilution
2 comprising from about 0.0005 to about 0.35% by weight of ketone acid and from about
3 0.0005 to about 0.15% by weight of iodine containing biocide, based upon 100% weight of
4 total composition.

1 34. The composition of claim 1, wherein the weight ratio of the aromatic
2 carboxylic acid to the iodine containing biocide ranges from about 0.0006:1 to about 1990:1.

1 35. The composition of claim 34, wherein the weight ratio of the aromatic
2 carboxylic acid to the iodine containing biocide ranges from about 0.0063:1 to about 1400:1.

1 36. The composition of claim 1, wherein said composition is a use dilution
2 comprising from about 0.00005 to about 0.4975% by weight of aromatic carboxylic acid and
3 from about 0.00005 to about 0.40% by weight of iodine containing biocide, based upon 100%
4 weight of total composition.

1 37. The composition of claim 36, wherein said composition is a use dilution
2 comprising from about 0.0005 to about 0.35% by weight of aromatic carboxylic acid and from
3 about 0.0005 to about 0.15% by weight of iodine containing biocide, based upon 100% weight
4 of total composition.

1 38. An antimicrobial composition comprising a synergistic mixture of:
2 (a) dehydroacetic acid or a salt thereof; and
3 (b) 3-iodo-2-propynyl butyl carbamate.

1 39. An antimicrobial composition comprising a synergistic mixture of:
2 (a) salicylic acid or a salt thereof; and
3 (b) 3-iodo-2-propynyl butyl carbamate.

1 40. An antimicrobial composition comprising a synergistic mixture of:
2 (a) dehydroacetic acid or a salt thereof;
3 (b) salicylic acid or a salt thereof; and
4 (c) 3-iodo-2-propynyl butyl carbamate.

- 1 41. A method of inhibiting the growth of microorganisms comprising applying an
- 2 effective amount of the composition of claim 1.

INTERNATIONAL SEARCH REPORT

Int ional Application No

PCT/US 02/06304

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A01N47/12 //A01N43/16,37/40

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, EPO-Internal, PAJ, BIOSIS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE WPI Derwent Publications Ltd., London, GB; AN 1992-102760 XP002207328 "Antimicrobial cream for cleaning optical lenses-contains 2-(4-thiazolyl)benzimidazole, oxy-bis:phenoxy-arsine, alpha-bromo:cinnamaldehyde..etc..as antimicrobial agent" & JP 04 049207 A (OLYMPUS OPTICAL CO LTD), 18 February 1992 (1992-02-18) ...	1,2,4-6, 9-11,13, 14, 17-23, 28,30, 31,34, 35,38-41
Y	abstract --- -/--	7,8

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

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- * & * document member of the same patent family

Date of the actual completion of the international search

25 July 2002

Date of mailing of the international search report

07/08/2002

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INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 02/06304

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE WPI Derwent Publications Ltd., London, GB; AN 1995-027903 XP002207329 "Antifungal waterproof cloths-has coating resin comprising 3-iodo-2-propynyl-N-N-butyl carbamate and e.g. 2-(4-thiazolyl)-benzimidazole" & JP 06 313269 A (OSAKA KASEI KK; TORAY IND INC), 8 November 1994 (1994-11-08)	1,2,4-6, 9-11,13, 14,17, 28-31, 38,41
Y	abstract	7,8
Y	US 5 906 981 A (GAGLANI KAMLESH D) 25 May 1999 (1999-05-25) column 1, line 6 - line 13 column 2, line 52 - line 61 column 10, line 21 - line 23	7,8

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 02/06304

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
JP 4049207	A	18-02-1992	JP 2634483 B2	23-07-1997
JP 6313269	A	08-11-1994	NONE	
US 5906981	A	25-05-1999	AT 211884 T	15-02-2002
			AU 3290197 A	05-01-1998
			BR 9710687 A	11-01-2000
			CA 2257372 A1	11-12-1997
			DE 69709659 D1	21-02-2002
			DK 926953 T3	01-07-2002
			EP 0926953 A1	07-07-1999
			WO 9746095 A1	11-12-1997

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